

AMPLIFICATION OF NUCLEIC ACIDS

CROSS-RELATED APPLICATIONS

5 This application is a continuation-in-part of U.S. Application No. 08/564,653, filed November 29, 1995.

TECHNICAL FIELD

10 The present invention relates generally to amplification of nucleic acids and, in particular, to amplification of G+C-rich templates.

BACKGROUND OF THE INVENTION

15 Amplification of nucleic acids has revolutionized molecular biology and is now one of the most rapid and versatile methods of producing large quantities of DNA for molecular analysis. Despite the power of this technique, uniform amplification of all templates, regardless of length or G+C content, remains a challenge. Amplification of longer nucleic acid fragments (up to 35 bp) has been accomplished by utilizing different enzymes selected for their thermal stability and editing functions (see *PCR Methods and Applications* 2:51, 257, 1993; Lundberg et al., *Gene* 108:1, 1991; 20 Mattila et al., *Nucleic Acids Res.* 19:4967, 1991; Barnes, *Proc. Natl. Acad. Sci. USA* 91:2216, 1994; Cheng et al., *Proc. Natl. Acad. Sci. USA* 91:5695, 1994; Jeffreys et al., *Nucleic Acids Res.* 16:10953, 1988; Krishnan et al., *Nucleic Acids Res.* 19:6177, 1991; Maga and Richardson, *Biotechniques* 11:185, 1991; Rychlik et al., *Nucleic Acids Res.* 18:6409, 1990; and Kainz et al., *Anal. Biochem.* 202:46, 1992). However, templates 25 with high G+C content, regardless of length, are only variably amplifiable or sometimes completely unamplifiable even in the presence of reagents that facilitate strand separation, stabilize the polymerase, or isostabilize DNA (e.g., dimethyl sulfoxide (DMSO), formamide, glycerol, or tetramethylammonium chloride (TMACl)) (Bookstein et al., *Nucleic Acids Res.* 18:1666, 1990; Sarkar et al., *Nucleic Acids Res.* 30 18:7465, 1990; Pomp and Medrano, *Biotechniques* 10:58, 1991; and Hung et al., *Nucleic Acids Res.* 18:4953, 1990).

 Although the average G+C content of the human genome is about 40%, individual genes and genetic elements may have G+C content that is higher. For example, the human *c-myc* gene is 60% G+C with regions of greater of 70% G+C. As 35 well, the majority of 5' ends of genes and promoters have regions that are G+C rich, and some diseases, such as Fragile X Syndrome, result from the *in vivo* expansion of G+C

rich triplets (e.g., CGG for Fragile X Syndrome). Detection of these triplets in both normal and diseased individuals is difficult. Furthermore, the inability to uniformly amplify DNAs with high G+C content hinders other methods, such as quantitation of transcripts, gene mapping and sequence analysis.

5 In the present invention, compositions and methods are provided for amplifying nucleic acids regardless of their G+C content as well as providing other related advantages.

SUMMARY OF THE INVENTION

10 This invention generally provides methods and compositions for increasing the efficiency of amplification of nucleic acids, especially nucleic acids with high G+C content.

 In one aspect, a reaction mixture is prepared by mixing nucleic acid templates, one or more primers, nucleotides, a first DNA polymerase and a second
15 DNA polymerase that has 3' exonuclease activity, and adding to the reaction mixture a zwitterion and a compound that disrupts base pairing in an amount sufficient to increase amplification of an 80% G+C, 500 bp DNA fragment by two-fold, when the zwitterion and the compound are present as compared to when the zwitterion and the compound are absent. In a related aspect, the reaction mixture is prepared by mixing a
20 homogeneous nucleic acid template, one or more primers, nucleotides, a first DNA polymerase and a second DNA polymerase that has 3' exonuclease activity, and adding to the reaction mixture a zwitterion or a compound that disrupts base pairings in an amount sufficient to increase amplification of an 80% G+C, 500 bp DNA fragment by two-fold, when the zwitterion or compound are present as compared to when the
25 zwitterion or compound are absent.

 In one embodiment, the first DNA polymerase lacks a 5' exonuclease activity. In a preferred embodiment, the first DNA polymerase is KlenTaq1 DNA polymerase and the second DNA polymerase is *Pfu* DNA polymerase. In other
30 embodiments, the DNA polymerase pairs are *rTth* DNA polymerase and *Thermococcus litoralis* DNA polymerase; *Taq* DNA polymerase and *Pyrococcus* DNA polymerase; *Taq* DNA polymerase and *Pwo* DNA polymerase.

 In other embodiments, the zwitterion is selected from the group consisting of betaine, monomethyl glycine, dimethyl glycine, and D-carnitine. In yet another embodiment, the compound that disrupts base pairing is dimethylsulfoxide or
35 formamide. In a preferred embodiment, the zwitterion is betaine and the compound is DMSO.

In yet other embodiments, the nucleic acid template is selected from the group consisting of genomic DNA, cDNA, plasmid DNA, DNA fragment, and viral DNA.

5 These and other aspects of the invention will become evident upon reference to the following detailed description and attached drawing. In addition, various references are set forth below which describe in more detail certain procedures or compositions. Each of these references are incorporated herein by reference in their entirety as if each were individually noted for incorporation.

10 BRIEF DESCRIPTION OF THE DRAWINGS AND SEQUENCE LISTING

Figures 1A and 1B are photomicrographs depicting the electrophoretic screening data for the various test fragments and test solutions according to the present invention.

15 Figure 2 is a photomicrograph depicting the electrophoretic screening data for the amplification of the CGG triplet repeat in Fragile X Syndrome.

Figure 3 is a photograph of amplification of cDNAs from a small number of cells and tissues. M, molecular weight markers; JTP, cDNA from activated Jurkat T cells; J1000, cDNA from 1000 Jurkat cells; J5000, cDNA from 5000 Jurkat cells; Ear1, cDNA from human inner ear; Ear2, cDNA from human inner ear.

20 SEQ ID No. 1 is a DNA sequence of the M13 forward primer.

SEQ ID No. 2 is a DNA sequence of the M13 reverse primer.

SEQ ID No. 3 is a DNA sequence of MHCcln1, which is derived from the MHC gene region.

SEQ ID No. 4 is a partial DNA sequence of human OCT-T1 cDNA.

25 SEQ ID No. 5 is a DNA sequence of human OCT1 cDNA.

SEQ ID No. 6 is a sequence of a primer used to amplify the repeat region of the Fragile X gene, FMR1.

SEQ ID No. 7 is a sequence of a primer used to amplify the repeat region of the Fragile X gene, FMR1.

30 SEQ ID No. 8 is a partial DNA sequence of TFR, the human transferrin receptor.

DETAILED DESCRIPTION OF THE INVENTION

35 Prior to setting forth the invention, it will be helpful to an understanding of the invention to define certain terms that are used hereinafter.

Nucleotides are depicted according to their recognized abbreviations, that is, "A" refers to adenine, "C" refers to cytosine, "G" refers to guanine, "T" refers to thymine, and "N" refers to either A or C or T or G.

As used herein, "amplification" refers to the increase in the number of
5 copies of a particular nucleic acid fragment resulting from an enzymatic reaction, such as polymerase chain reaction, ligase chain reaction, or the like.

As used herein, "oligonucleotide" refers to a nucleic acid molecule comprising two or more deoxyribonucleotides or ribonucleotides, and preferably more than three. An oligonucleotide may be synthesized or produced by amplification or
10 cloning.

As used herein, "polymerase chain reaction" or "PCR" refers to a particular process of amplification for the exponential amplification of a specific DNA fragment by utilizing two oligonucleotide primers that hybridize to opposite strands and flank the region of interest in a target DNA. The process consists of a repetitive series
15 of cycles involving template denaturation, primer annealing, and the extension of the annealed primers by a DNA polymerase, such as the thermostable *Taq* DNA polymerase.

As used herein, a "primer" is an oligonucleotide that anneals to one strand of a nucleic acid template and allows elongation of a complementary strand by a
20 polymerase.

As used herein, "template" refers to a sequence of nucleotides that can be copied by a DNA polymerase from a primer bound to the template.

As used herein, a "zwitterion" refers to a molecule that is a dipolar ion. At the isoelectric pI of the molecule, it will have no net charge; at a pH range between
25 approximately the K_a and the K_b , the molecule bears little net charge.

I. Amplification of nucleic acid templates

Amplification using thermostable enzymes generally involves multiple cycles of denaturation, annealing, and synthesis performed at different temperatures.
30 An amplification reaction typically contains either heterogeneous or homogenous template nucleic acid, one or more oligonucleotide primers, nucleotides, various buffers and cofactors, and DNA polymerase. To amplify heterogeneous G+C rich templates, a zwitterion and compound that disrupts base pairing are added to the mix; if the template is homogeneous (*e.g.*, plasmid clone DNA), either a zwitterion or compound is added.

35 Protocols vary somewhat in concentrations of reagents, temperatures, and incubation times, depending upon the apparatus, length of amplified product,

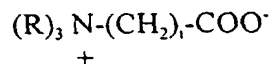
enzyme used, and other parameters. Protocols are readily available (*see, for example*, manufacturer's instructions, Ausubel et al., *Current Protocols in Molecular Biology*, Greene Publishing, 1995; and Examples herein). Generally, a "hot-start" procedure is used, so that all reagents, save one, such as nucleotides, are mixed, and the temperature is raised to at least annealing temperature before the saved reagent is added. Amplification reactions may be analyzed by any of a variety of known methods. For example, the reaction may be electrophoresed in a polyacrylamide or agarose gel. The amplified material may be detected by staining with a dye, such as ethidium bromide, or by autoradiography if a radioactive label is incorporated.

II. Zwitterions and compounds that disrupt base pairing

As noted above, the present invention provides compositions comprising a zwitterion or dipolar ion in combination with a compound that disrupts base pairing to enhance and improve amplification of nucleic acids that are G+C rich. Typically, examples of G+C rich regions that show improvement of amplification in the present invention contain about 60% G+C over greater than 50 nucleotides, about 80% G+C in up to 500 nucleotide region, and 100% G+C in up to about 200 nucleotides.

A zwitterion contains both an acidic group and a basic group. The acidic group is associated and the basic group is dissociated at pH values between about the pK_a (the pH at which the acidic group is half associated and half dissociated) and about the pK_b (the pH at which the basic group is half associated and half dissociated). Dipolar ions are encountered, for example, whenever a molecule contains both an amino group and an acid group, providing that the amine is more basic than the anion of the acid. For example, all amino acids exist in a zwitterionic form between about pH 4 and about pH 9. Over this pH range, the amino acids bear little net charge, and at the isoelectric point, pI , bear no net charge.

The zwitterions for use in the present invention must have a basic group containing at least one alkyl group, which is preferably a methyl group. Preferably the basic group is an amine. A general formula for the zwitterion is thus:



wherein R is H, CH_3 , CH_2-CH_3 , or $(CH_2)_j-CH_3$, at least one R is not H, i is from 2 to about 6, and j is from 2 to about 6. Preferred zwitterions are betaine (trimethyl glycine), D-carnitine, dimethyl glycine, and monomethyl glycine.

For use in the present invention, the zwitterion should not inhibit DNA polymerases. It is preferable that the DNA polymerase retain at least 90% of activity at 2.5 M zwitterion concentration. The effect of the zwitterion on nucleic acid molecules is a decrease in the T_m of GC base pairs and an increase in the T_m of AT base pairs. The decrease or increase is preferably at least 2°C, at least 3°C, at least 4°C, and most preferably at least 5°C.

Compounds that disrupt base pairing (e.g., disrupt hydrogen bonding) include DMSO, formamide, sodium perchlorate, glyoxyl, and the like. Such compounds may be identified by their ability to lower the melting temperature of a nucleic acid duplex. Such compounds should not inhibit DNA polymerases. It is preferable that the DNA polymerase retain at least 90% of activity at a 10% compound concentration.

A zwitterion or a compound that disrupts base pairing can be tested for utility in the context of the present invention by demonstration of at least a two fold increase in the amount of amplified product from a homogeneous template of at least 80% G+C, in the presence of the zwitterion or compound as compared to the absence of the zwitterion or compound. When heterogeneous templates (i.e., multiple templates) are to be amplified, the zwitterion and compound that disrupts base pairing are tested in combination. The zwitterion and compound are added in amounts sufficient to cause an increase in efficiency of amplification. In general, the zwitterion is added from about 0.1 M to about 3 M, from about 0.5 M to about 4.5 M, or from about 1 M to about 2.5 M. In general, the compound is added from about 5% to about 15%, or preferably from about 5% to about 10%.

III. Two enzyme amplification systems

As noted above, an improvement in amplification is achieved when amplification is performed in a two-enzyme system.

Two-enzyme systems are used for amplification of long fragments. In general, one of the enzymes has a proofreading activity ($3' \rightarrow 5'$ exonuclease) that corrects nucleotide misincorporations that may otherwise prematurely terminate DNA synthesis, and the other enzyme is highly processive. Preferably, one or both of the enzymes lack $5' \rightarrow 3'$ exonuclease activity. Also, preferably, the enzymes are thermostable. Examples of combinations of enzymes that are commonly used are KlenTaq1 plus *Pfu* DNA polymerases (1:15 v/v), *rTth* plus Vent® isolated from *Thermococcus litoralis* DNA polymerases (Perkin-Elmer, Foster City, CA); *Taq* and *Pyrococcus* DNA polymerases (Life Technologies, Gaithersburg, MD); and *Taq* and

Pwo DNA polymerases (Boehringer-Mannheim, Indianapolis, IN). Other combinations may be used.

IV. Application of amplification of G+C rich templates

5 As described herein, the present invention provides compositions and methods for uniform amplification of nucleic acid templates regardless of G+C content. Such amplification allows accurate quantitation, detection of G+C rich regions, diagnosis of certain diseases, and improves construction of cDNA libraries, DNA sequence analysis, and the like. For example, 5' ends of genes are G+C rich and are
10 often not amplified well during construction of a cDNA library. Some coding sequences are G+C rich and will also be underrepresented in a cDNA or genomic library. Without the present invention, if the starting material was a cDNA library with G+C-rich cDNA present in medium abundance (<0.1%), upon amplification, this G+C-rich nucleic acid would be less efficiently amplified and gradually lost. Thus, to find
15 the cDNA in a screen by hybridization, several million clones would likely need to be examined. Using the present invention, the G+C rich cDNA should be present in the amplified material at the same level as in the starting material. Following amplification of cDNA from RNA isolated from a small number of cells, differential display amplification can be performed.

20 Moreover, several genetic diseases are caused by a natural expansion of a G+C-rich triplet. Such diseases include fragile X syndrome, spinobulbar muscular atrophy, myotonic dystrophy, Huntington's chorea, and spinocerebellar ataxia type 1.

Furthermore, in addition to enhancing amplification, the present invention may improve the efficiency of nucleic acid synthesis in general, including
25 reverse transcription, *in vitro* transcription, primer extension and the like.

EXAMPLES

EXAMPLE 1

30 AMPLIFICATION OF G+C RICH TEMPLATES USING *Taq* DNA POLYMERASE
AND BETAINE OR DMSO

A number of test DNAs are utilized in amplification reactions. Four of the test fragments are:

(i) a 2.7 kb cDNA encoding the octamer binding protein, OCT1 (GenBank accession No. X13403; Sturm et al., *Genes & Dev.* 2:1582, 1988; SEQ ID No. 8), which has a 52% G+C content;

(ii) a 1.6 kb fragment from the MHC region, MHCc1n1, which has a 64% G+C content (see Shukla et al., *Nucleic Acids Res.* 19:4233, 1991; GenBank Accession No. L20433; SEQ ID No. 3, which presents nucleotides 320 to 850 of L20433);

(iii) a 0.8 kb fragment of human OCT-T1 cDNA, which encodes a transcription factor expressed in T lymphocytes that binds the octamer sequence, and which has an 80% G+C content (see Bhargava et al., *Proc. Natl. Acad. Sci. USA* 91:10260, 1993; SEQ ID NO. 4); and

(iv) a 0.72 kb fragment of human transferrin receptor, TFR, which has a 44% G+C content (*Nature* 311:675, 1984; GenBank Accession No. X01060; SEQ ID No. 5, which presents nucleotides 352 to 866 of X01060).

As each of these fragments are cloned in pBluescript or a pUC plasmid (MHCc1n1), primers for amplification of these fragments are the M13 forward primer (CGCCAGGGTTTCCAGTCACGAC ; SEQ. ID No. 1) and the M13 reverse primer (AGCGGATAACAATTCACACAGGA ; SEQ ID No. 2). Thus, the sizes of the amplified fragments for each of these four templates, with the exception of MHCc1n1, include 200 bp of multiple cloning sites of pBluescript vector.

The two G+C-rich templates are unsuccessfully amplified (MHCc1n1 and OCT-T1) using AmpliTaq® DNA polymerase (Perkin-Elmer Cetus) under standard conditions. (See *Science* 239:487, 1988). The addition of DMSO (10% v/v), betaine (up to 2.5 M), glycerol (10% v/v), formamide (10% v/v) and TMACl (tetramethylammonium chloride) (0.1 mM to 1.0 M) or increasing annealing and denaturation temperatures did not result in visible amplification. In addition, betaine (>1.0 M), formamide (>5%), and TMACl (>100 mM) are inhibitory for amplification of the non-G+C-rich template OCT1.

EXAMPLE 2

AMPLIFICATION OF G+C RICH TEMPLATES USING A TWO ENZYME DNA POLYMERASE SYSTEM AND BETAINE OR DMSO

Each of the G+C-rich templates is set up in an amplification reaction using a two enzyme mixture, with and without the addition of betaine (0.5 to 2.5 M),

DMSO (5-10%), formamide (5-10%), glycerol (5-10%), TMACl (0.1 mM to 1 M), and tetramethylammonium acetate (TMAA) (0.1 mM - 1.0 M).

The amplification reaction (50 μ l) is performed in 20 mM Tris-HCl, pH 9.0 at 25°C, containing 150 μ g/ml of bovine serum albumin, 16 mM ammonium sulfate, 2.5 mM magnesium chloride, 200 μ M of each dNTP, 50 ng each of the M13 forward primer and the M13 reverse primer, 0.5-1.0 ng of plasmid templates and 0.2-0.4 μ l of Klentaq LA16 (a 1:15 v/v ratio of Klentaq1 (Ab Peptides Inc., St. Louis, MO), which is *Taq* DNA polymerase lacking the 5' to 3' exonuclease activity, and *Pfu* DNA polymerase (Stratagene, LaJolla, CA); the long and accurate (LA) system described in Barnes, *Proc. Natl. Acad. Sci. USA* 91:2216, 1994). DMSO, TMACl, TMAA, betaine, or the combination of DMSO and betaine is added to aliquots of the same mixture without dNTPs and overlaid with mineral oil. Amplification is performed in a thermocycler instrument using thin-walled tubes (0.5 ml). The tubes are heated at 95°C for 15 seconds and amplification is initiated by the addition of dNTPs at 80°C.

The following cycling conditions are used for 25 cycles: denaturation at 94°C for 15 seconds, annealing at 50 to 55°C for 1 minute, and extension at 68°C for 5 minutes. Following amplification, 10 μ l of the resulting product is loaded onto a 1% agarose gel (see *Molecular Cloning: A Laboratory Manual* by J. Sambrook et al., Cold Spring Harbor, 1989). The amplified DNA is stained with ethidium bromide and the gel is photographed while UV-irradiated by a transilluminator.

The results are shown in Figures 1A and 1B. The individual numbered lanes in Figures 1A and 1B, contain the template and additives listed in the following table.

TABLE 1

Lane	Template*	Additive	Lane	Template	Additive
1	OCT1	2.2 M betaine	10	all four	10% DMSO
2	OCT1	10% DMSO	11	all four	1.1 M betaine
3	TFR	2.2 M betaine	12	all four	2.2 M betaine
4	TFR	10% DMSO	13	all four	5% DMSO + 1.1 M betaine
5	MHCcIn1	2.2 M betaine	14	all four	5% DMSO + 2.2 M betaine
6	MHCcIn1	10% DMSO	15	all four	10% DMSO + 1.1 M betaine
7	OCT-T1	2.2 M betaine	16	all four	10% DMSO + 2.2 M betaine
8	OCT-T1	10% DMSO	17	all four	7% DMSO + 1.0 M betaine
9	all four	5% DMSO	18	all four	5% DMSO + 1.4 M betaine

* "all four" refers to the mixture of the 4 templates, OCT1, OCT-T1, TFR and MHCclnl.

As shown in Figure 1A, each of the templates could be amplified in the presence of betaine or DMSO. Furthermore, DMSO and betaine do not have a negative effect on amplification. In contrast, the addition of tetramethylammonium chloride (TMACl), which is also a tetraalkylammonium ion, does not have a similar effect as that found with betaine. Moreover, TMACl and TMAA (at 0.1-1.0 M) inhibit amplification of OCT1. The addition of betaine and/or DMSO aids in the amplification of G+C-rich templates (lanes 5-8, Figure 1A), while not having any adverse effect on the amplification of the other templates (lanes 1-4, Figure 1A).

Amplification is also performed on a mixture of the four templates, varying the concentrations of DMSO and/or betaine. As can be seen in Figure 1B, the four templates are not amplified uniformly upon addition of either DMSO or betaine alone (lanes 9-12). All the templates were amplified to a varying extent in the presence of 2.2 M betaine (lane 12), 5% DMSO + 2.2 M betaine (lane 14), or 10% DMSO + 1.1 M betaine (lane 5). When 7% DMSO with 1.0 M betaine and 5% DMSO in combination with 1.4 M betaine (lanes 17 and 18) were tested, all templates were amplified uniformly. Thus, by adding both DMSO and betaine, preferably within a range of 1.0 M betaine with 6-8% DMSO or 5% DMSO with 1.2-1.8 M betaine, all four templates are amplified.

Thus, approximately equimolar amplification of all the templates occurred at 1.0 M betaine + 7% DMSO (lane 17), and 5% DMSO + 1.4 M betaine (lane 18). A combination of 1.0 M betaine with 6-8% DMSO or 5% DMSO with 1.2-1.8 M betaine resulted in similar uniform amplifications; accordingly such concentrations are preferred. Although at a higher concentration of DMSO (12-15%) or betaine (1.8-2.5 M), all the templates showed variable amplification, approximately equimolar yield of individual components of the template mixtures were obtained when DMSO and betaine were within the preferred concentration range and were combined with the LA system.

EXAMPLE 3AMPLIFICATION OF TEMPLATES USING TWO ENZYME DNA POLYMERASE SYSTEMS AND
BETAINE OR DMSO

- 5 Various single and two enzyme systems were used to amplify OCT-T1 (80% G+C) or TFR (44% G+C) templates. Amplification reactions were performed as above in the presence of 2 M betaine. The results are presented below.

Table 2

Enzyme System	Enzymes	OCT-T1 (80% G+C)	TFR (44% G+C)	Sensitivity to Betaine
LA-16	Klentaql and <i>Pfu</i> DNA Polymerase	+++	+++	-
AmpliTaq® DNA Polymerase	<i>rTaq</i> DNA Polymerase	-/+	+++	+
XL PCR (Perkin-Elmer)	<i>rTth</i> and <i>T. litoralis</i> DNA Polymerase	+++	+++	-
Elongase (BRL)	<i>Taq</i> and <i>Pyrococcus</i> DNA Polymerase	+++	+++	-
Expand System (Boehringer Mannheim)	<i>Taq</i> and <i>Pwo</i> DNA Polymerase	++	+++	-
Vent® DNA Polymerase (NE Biolabs)	<i>T. litoralis</i> Polymerase	-	+++	+
Deep Vent™ Polymerase (NE Biolabs)	<i>Pyrococcus</i> Polymerase	-	-	+

10

As shown above, all enzymes, except *Pyrococcus* polymerase, amplified TFR, a template with average G+C content. However, only the two enzyme systems were able to amplify OCT-T1, a template with high G+C content.

EXAMPLE 4

AMPLIFICATION OF TEMPLATES USING STRUCTURAL ANALOGUES OF BETAINE

Various analogues of betaine and two tetraalkylammonium salts are used to amplify OCT-T1 or TFR templates. Amplification reactions are performed as above in the presence of either 2 M analogue or at least 0.15 M salt. Results are presented in the Table below.

Table 3

Compound	OCT-T1	TFR
Monomethyl glycine (2 M)	-/+	+++
Dimethyl glycine (2 M)	+	+++
Glycine betaine (2 M)	+++	+++
D-carnitine (2 M)	+++	+++
TMAC (>0.15 M)		-
TMAA (>0.15 M)	-	-

10

The structure of the compounds is shown below.

Table 4

Glycine betaine	$(\text{CH}_3)_3\text{N}^+-\text{CH}_2-\text{COO}^-$
D-carnitine	$\text{OOC}-\text{CH}_2-\overset{\text{OH}}{\underset{ }{\text{CH}}}-\text{CH}_2-\text{N}^+(\text{CH}_3)_3$
Monomethyl glycine	$\text{CH}_3-\text{H}_2\text{N}^+-\text{CH}_2-\text{COO}^-$
Dimethyl glycine	$(\text{CH}_3)_2-\text{NH}^+-\text{CH}_2-\text{COO}^-$
TMACl	$(\text{CH}_3)_4-\text{N}^+-\text{Cl}^-$
TMAA	$(\text{CH}_3)_4-\text{N}^+-\text{CH}_2\text{COO}^-$

As shown in the table, TMACl and TMAA are unable to facilitate amplification of either template and most likely inhibit the enzyme. However, the

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zwitterions facilitated amplification of both templates, the quarternary ammonium ions working best.

EXAMPLE 5

5 AMPLIFICATION OF G+C RICH REPEATS OF THE FRAGILE X GENE

Amplification of the G+C-rich repeat expansion involved in Fragile X Syndrome is performed. Fragile X Syndrome is one of the most common causes of mental retardation affecting 1 in 1,500 males, and 1 in 2,500 females. The fragile X site has been mapped to Xq27.3, and shown to be due to the expansion of CGG triplet repeats in the 5'-end of the FMR1 gene. Normal subjects have 2-50 CGG repeats with an average at 29 repeats, those with premutation have about 50-200 repeats, and individuals with disease have more than 200 repeats (*i.e.*, a full mutation) (see Nussbaum and Ledbetter, *Ann. Rev. Genetics* 20:109, 1986; Verkerk et al., *Cell* 65:905, 1991; and Kremer et al., *Science* 252:1711, 1991). Diagnostic confirmation of this disease has primarily been accomplished by cytogenetic analysis of the cells and Southern blotting of the DNA from these patients (see Sutherland, *Am. J. Hum. Genet.* 31:125, 1979; and Rousseau et al., *N. Engl. J. Med.* 325:1673, 1991). The sequence of the fragile X CGG repeat region may be found in GenBank accession No. X61378.gb-pr, and encompasses nucleotides 2599 to 2868.

The methods described herein are used to amplify the G+C-rich fragile X CGG triplet repeat region (85% G+C) directly from genomic DNA of a cell line with a normal FMR1 gene, JV, (29 CGG repeat), genomic DNA from 2 males with premutations (74 and 119 CGG repeats, respectively), and genomic DNA from 2 patients carrying full mutations (>1000 CGG repeats). The sizes of the CGG repeats were obtained by Southern blotting and hybridization with specific probes.

Amplification is performed as described above in the presence of 10% DMSO + 1.1 M betaine or 5% DMSO + 2.0 M betaine. The sequence of primers used are:

30 primer c: GCTCAGCTCC GTTTCGGTTT CACTTCCGGT (SEQ ID No. 6)
 primer f: AGCCCCGCAC TTCCACCACC AGCTCCTCCA (SEQ ID No. 7)

35 Either 10 or 100 ng of genomic DNA from each sample is mixed in 20 mM Tris-Cl (pH 9.0) containing 150 µg/ml bovine serum albumin, 16 mM ammonium sulfate, 2.5 mM magnesium chloride, and 50 ng of each primer. All the

components, except dNTP, are denatured at 94°C for 15 sec, and dNTPs are added at 80°C. Amplification proceeded using the following cycling conditions for 40 cycles: denaturation at 94°C for 15 sec, annealing at 65°C for 1 minute, and extension at 68°C for 5 min (or annealing and extension was alternatively done at 68°C for 5 min).

- 5 Following amplification, 10 µl of the reaction was loaded on a 2% agarose gel and the bands visualized by ethidium bromide staining.

The results are depicted in Figure 2, in which lane 1 is JY DNA (10 ng); lane 2 is JY DNA (100 ng); lane 3 is 74 CGG repeat containing DNA; lane 4 is 119 CGG repeats containing DNA; lane 5 is DNA containing heterozygous full mutation in a female patient; and lane 6 is a full mutation from a male DNA sample.

10 An approximately 300 bp PCR product is observed for JY (Figure 2, lanes 1 and 2), and the expected size fragments for 74 repeat (440 bp) and 119 repeats (575 bp) are observed as well (Figure 2, lanes 3 and 4). However, no specific band is observed for the DNA from a male patient carrying a full mutation (Figure 2, lane 5), and only a normal band (300 bp) is observed for DNA from a female patient carrying a full mutation as well as a normal gene (Figure 2, lane 6). Thus, the normal sized repeat (with an average repeat number of 29) and remutation expansion can be visibly amplified, whereas the full mutation (>200 repeat) is not visibly amplified.

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EXAMPLE 6

AMPLIFICATION OF cDNAs USING BETAINE OR DMSO

First strand cDNAs synthesized from total RNA isolated from a small number of cells and tissues are amplified. Either Jurkat T cells, induced for 4 hrs with 50 nM phorbol 12-myristate 13-acetate and 2 µg/ml phytohemagglutinin, or human inner ear samples, obtained during a surgical procedure, were used. Total RNA was isolated, and double-stranded cDNA was made. An adaptor containing a *NotI* site was ligated to the cDNAs. Amplification reactions are performed as described above using Klentaq LA16, using the adaptor as primer in the presence of 1.5 M betaine and 7% DMSO, except that each cycle consisted of denaturation at 95°C for 15 sec, annealing at 55°C for 1 min, and extension at 68°C for 5 min. A second round of amplification was performed on 1 µl of reaction mix using the cycle conditions: denaturation at 95°C for 15 sec, annealing at 60°C for 1 min, and extension at 68°C for 5 min.

35 As shown in Figure 3, uniform amplification occurred using cDNA from tissues (Ear1 and Ear2) as well as from cell lines (JTP, J1000, J5000). Moreover, even

It will be appreciated that, although specific embodiments of the invention have been described herein for purposes of illustration, various modifications may be made without departing from the spirit and scope of the invention. Accordingly, the invention is not limited except as by the appended claims.

SEQUENCE LISTING

(1) GENERAL INFORMATION:

(i) APPLICANTS: Sherman M Weissman
Namadev Baskaran

(ii) TITLE OF INVENTION: AMPLIFICATION OF NUCLEIC ACIDS

(iii) NUMBER OF SEQUENCES: 8

(iv) CORRESPONDENCE ADDRESS.

(A) ADDRESSEE SEED and BERRY LLP
(B) STREET: 701 Fifth Avenue, 6300 Columbia Center
(C) CITY: Seattle
(D) STATE: Washington
(E) COUNTRY: USA
(F) ZIP: 98104

(v) COMPUTER READABLE FORM.

(A) MEDIUM TYPE: Floppy disk
(B) COMPUTER: IBM PC compatible
(C) OPERATING SYSTEM: PC-DOS/MS-DOS
(D) SOFTWARE: PatentIn Release #1 0. Version #1.30

(vi) CURRENT APPLICATION DATA:

(A) APPLICATION NUMBER:
(B) FILING DATE: 29-NOV-1996
(C) CLASSIFICATION

(vii) ATTORNEY/AGENT INFORMATION:

(A) NAME: Nottenburg Ph D., Carol
(B) REGISTRATION NUMBER: 39.317
(C) REFERENCE/DOCKET NUMBER: 390036.402C1

(ix) TELECOMMUNICATION INFORMATION:

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(2) INFORMATION FOR SEQ ID NO.1.

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 24 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION. SEQ ID NO:1:

GGATCCGGCC	CCACGGAGGT	CCCCATCTCC	CTCAAGATTC	TCAGATTCAT	CCCCAATGAG	60
TGGTGTAGCC	CCTACAGGGG	TGTCAGCCCC	CCTCATCACC	AACAGTGACA	GTGACAGAGG	120
CTGGAGATGA	GGGGCCAGCA	GGCTCCAGGG	AGTCGGGGGT	GGCCTTGGGC	AGGGTTTCTT	180
CACTACGAGG	GGTGTTC CCC	AAAGAGCCAT	GAAGTGTAGA	GGAAGAGAAA	AAGTTCAGAG	240
CTAAGGGCTC	AGGAGATCCT	GTGTATTTAG	GGAAGGTGAC	GGTCCAATTG	GGGCCCCGTT	300
TAGCTGCACT	CACCTCTCTC	GGTGGCTCCT	CTGGTTTCCT	TCTCCAGCAG	CTCCCCCATC	360
TCAGCGGGGG	CCATCCCCCT	GGGAGGGGAG	ACAAGGGACA	GGAGGGCTGG	TCAGCCCAGT	420
AGAGAGTTGG	GGGGTCCAGG	ATGCCTGGGC	CCTGGGAAGA	GAGAGTAGGC	TCCGGGGCCT	480
ACCTCTTCCT	CTGGCCCTTC	CGCGGCCTCG	GCTGCCCGGA	GCCGCACAAC	CCTCCCCGGG	540
CCGCATAATC	CCTCCTTGAT	GACCTCCCT	CTCGGTAGTA	CCCGCACTCT	GGGGCCGAGA	600
GAAGAGGAGG	GGGCACGGAC	TCTTGGGGGG	GGCCTCCGAG	CCCGGCCCCG	CCCCTCTCCC	660
CGGCTGCACG	CGCCGATACC	CTTTGTACCC	AGGCGCGGGA	CCCGGACAAT	CCTCAGATCC	720

TCCAGCACCC GCTGCCCCC AGCCCGGTGG ACGGCCCTC GTGCCCTCA CGCGTGCTCC	780
TGGGGCCCCG GCGCCGTCG CCCAGTGGG GCAGGCCGGC GGCTGCACGC GCGCTCCGT	840
GCCCACTCCC CCCACCTCCC ACACCCTGGT CCCCTCATCC GCCCCGGTG CTGGCCCCCT	900
GGATTGCTGC AAGTCCCGCC CGGCCCCCGG CCCCGTTGCA CCCCCGAGC ATTGCACGGC	960
GCTTCCCCCG GGGGCGCGG CGGGCATGCA CCCGCCTCTC CCCCTCCCTT CCGCACCTCG	1020
GCGGCCGCCG CCGCTGCAGC TCCGCGGCC GCCGCCATCG CGCTTGCCT GGGGGCCGAG	1080
CCGGCGCGCG GCCGCCCCG GTCACGTGGG CGAGGGAGGG AGGGCGAGGA GGAGCCTTAA	1140
AGGAGCCGCT ACATGCTTTT TGGCCATTTT CCCCTGAGAG CGGCCTCGGA GATGGCTGTG	1200
ACTGTCCTAA GCTGGGAGCT GCAAGGGAGA ATTCCTGTCA TTCCTGGCCT CAGTTCTGCA	1260
GGGACCGAGG GCGAGACACG CCTGGGCCCA GGTGTGGCGT CTCTGTCCCC ATCTGGTTTT	1320
AGGTAACAAG CGGACGTTCT GAACTTCTCG GCTCTCGGCA GCGGCTGTAT TTCCTCTGGC	1380
CTGGTTGGGC TTTTCCCGCC TCTGGTTGCT TTTCTGCCTT TCTAGTTTTT GGGTTACCAG	1440
ATAGAAGGCT TGGCCTCAGT TTTGGCCTCG CCTTTTGTCT CTTTCTAACG AGCACGAAGG	1500
GGCGATAGGG ACGCGGAGGA CACCTTTATT CTTGGCTGGT TCTAGCATGC TGCTTCATGT	1560
CCCCTGGAGC AGCGTGCCCT TCTGAAAACC TGTGGCTAAA TGTCTCTTCT GTTTATATCT	1620
GGCGTGTTAC ACCTTCACAC GCACTAGGAT CC	1652

(2) INFORMATION FOR SEQ ID NO:4:

- (1) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 530 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(x1) SEQUENCE DESCRIPTION: SEQ ID NO:4:

CGAGGCCATC CGGCGGGCCT GCCTGCCAC GCCCGCGTG CAGAGCAACC TCTTCGCCAG	60
CCTGGACGAG ACGCTGCTGG CGCGGGCCGA GGCCTGGCG GCCGTGGACA TCGCCGTGTC	120
CCAGGGCAAG AGCCATCCTT TCAAGCCGGA CGCCACGTAC CACACGATGA ACAGCGTGCC	180
GTGCACGTCC ACTTCCACGG TGCTCTGGC GCACCACCAC CACCACCACC ACCACCACCA	240

GGCGCTCGAA CCCGGCGATC TGCTGGACCA CATCTCCTCG CCGTCGCTCG CGCTCATGGC 300
 CGGCGCGGGC GGC CGCGGGCG CGGCGGCCGG CGGCGGCGGC GCCCAGGACG GCCCGGGGGG 360
 CGGTGGCGGC CCGGGCGGCG GCGGCGGCC GGGCGGCGGC GGCCCCGGG GAGGCGGCGG 420
 TGGCGGCCCG GGGGGCGGCG GCGGCGGCC GGGCGGCGGC CTCCTGGGCG GCTCCGCGCA 480
 CCCTCACCCG CATATGCACA GCCTGGGCCA CCTGTGCAC CCCGCGGCGG 530

(2) INFORMATION FOR SEQ ID NO:5:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 515 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(x1) SEQUENCE DESCRIPTION: SEQ ID NO:5:

ATGGCGATAA CAGTCATGTG GAGATGAAAC TTGCTGTAGA TGAAGAAGAA AATGCTGACA 60
 ATAACACAAA GGCCAATGTC ACAAACCAA AAAGGTGTAG TGGAAGTATC TGCTATGGGA 120
 CTATTGCTGT GATCGTCTTT TTCTTGATTG GATTTATGAT TGGCTACTTG GGCTATTGTA 180
 AAGGGGTAGA ACCAAAACT GAGTGTGAGA GACTGGCAGG AACCGAGTCT CCAGTGAGGG 240
 AGGAGCCAGG AGAGGACTTC CCTGCAGCAC GTCGCTTATA TTGGGATGAC CTGAAGAGAA 300
 AGTTGTCGGA GAACTGGAC AGCACAGACT TCACCAGCAC CATCAAGCTG CTGAATGAAA 360
 ATTCATATGT CCCTCGTGAG GCTGGATCTC AAAAAGATGA AAATCTTGGC TTGTATGTTG 420
 AAAATCAATT TCGTGAATTT AACTCAGCA AAGTCTGGCG TGATCAACAT TTTGTTAAGA 480
 TTCAGGTCAA AGACAGCGCT CAAAACTCGG TGATC 515

(2) INFORMATION FOR SEQ ID NO:6:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 30 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

GCTCAGCTCC GTTTCGGTTT CACTTCCGGT

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(2) INFORMATION FOR SEQ ID NO:7:

- (1) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 30 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(x1) SEQUENCE DESCRIPTION: SEQ ID NO:7:

AGCCCCGCAC TTCCACCACC AGCTCCTCCA

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(2) INFORMATION FOR SEQ ID NO:8:

- (1) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 2584 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(x1) SEQUENCE DESCRIPTION: SEQ ID NO:8:

GAGGAGCAGC GAGTCAAGAT GAGAGTTCAG CCGCGGCGGC AGCAGCAGCA GACTCAAGAA	60
TGAACAATCC GTCAGAAACC AGTAAACCAT CTATGGAGAG TGGAGATGGC AACACAGGCA	120
CACAAACCAA TGGTCTGGAC TTTCAGAAGC AGCCTGTGCC TGTAGGAGGA GCAATCTCAA	180
CAGCCCAGGC GCAGGCTTTC CTTGGACATC TCCATCAGGT CCAACTCGCT GGAACAAGTT	240
TACAGGCTGC TGCTCAGTCT TTAAATGTAC AGTCTAAATC TAATGAAGAA TCGGGGGATT	300
CGCAGCAGCC AAGCCAGCCT TCCCAGCAGC CTTCACTGCA GGCAGCCATT CCCCAGACCC	360
AGCTTATGCT AGCTGGAGGA CAGATAACTG GGCTTACTTT GACGCCTGCC CAGCAACAGT	420
TACTACTCCA GCAGGCACAG GCACAGGCAC AGCTGCTGGC TGCTGCAGTG CAGCAGCACT	480
CCGCCAGCCA GCAGCACAGT GCTGCTGGAG CCACCATCTC CGCCTCTGCT GCCACGCCCA	540
TGACGCAGAT CCCCTGTCT CAGCCCATA CAGATCGACA GGATCTTCAA CAACTGCAAC	600
AGCTTCAACA GCAGAATCTC AACCTGCAAC AGTTTGTGTT GGTGCATCCA ACCACCAATT	660

TGCAGCCAGC GCAGTTTATC ATCTCACAGA CGCCCCAGGG CCAGCAGGGT CTCCTGCAAG	720
CGCAAAATCT TCAAACGCAA CTACCTCAGC AAAGCCAAGC CAACCTCCTA CAGTCGCAGC	780
CAAGCATCAC CCTCACCTCC CAGCCAGCAA CCCCAACACG CACAATAGCA GCAACCCCAA	840
TTCAGACACT TCCACAGAGC CAGTCAACAC CAAAGCGAAT TGATACTCCC AGCTTGAGAG	900
AGCCCAGTGA CCTTGAGGAG CTTGAGCAGT TTGCCAAGAC CTTCAAACAA AGACGAATCA	960
AACTTGATT CACTCAGGGT GATGTTGGGC TCGCTATGGG GAACTATAT GGAAATGACT	1020
TCAGCCAAAC TACCATCTCT CGATTTGAAG CCTTGAACCT CAGCTTTAAG AACATGTGCA	1080
AGTTGAAGCC ACTTTTAGAG AAGTGGCTAA ATGATGCAGA GAACCTCTCA TCTGATTCGT	1140
CCCTCTCCAG CCCAAGTGCC CTGAATTCTC CAGGAATTGA GGGCTTGAGC CGTAGGAGGA	1200
AGAAACGCAC CAGCATAGAG ACCAACATCC GTGTGGCCTT AGAGAAGAGT TTCTTGAGAG	1260
ATCAAAAGCC TACCTCGGAA GAGATCACTA TGATTGCTGA TCAGCTCAAT ATGGAAAAAG	1320
AGGTGATTCG TGTTTGTTTC TGTAACCGCC GCCAGAAAGA AAAAAGAATC AACCCACCAA	1380
GCAGTGGTGG GACCAGCAGC TCACCTATTA AAGCAATTTT CCCAGCCCA ACTTCACTGG	1440
TGGCGACCAC ACCAAGCCTT GTGACTAGCA GTGCAGCAAC TACCCTCACA GTCAGCCCTG	1500
TCCTCCCTCT GACCAGTGCT GCTGTGACGA ATCTTTCAGT TACAGGCACT TCAGACACCA	1560
CCTCCAACAA CACAGCAACC GTGATTTCCT CAGCGCCTCC AGCTTCCTCA GCAGTCACGT	1620
CCCCCTCTCT GAGTCCCTCC CTTTCTGCCT CAGCCTCCAC CTCCGAGGCA TCCAGTGCCA	1680
GTGAGACCAG CACAACACAG ACCACCTCCA CTCCTTTGTC CTCCCCTCTT GGGACCAGCC	1740
AGGTGATGGT GACAGCATCA GGTTTGCAAA CAGCAGCAGC TGCTGCCCTT CAAGGAGCTG	1800
CACAGTTGCC AGCAAATGCC AGTCTTGCTG CCATGGCAGC TGCTGCAGGA CTAAACCCAA	1860
GCCTGATGGC ACCCTCACAG TTTGCGGCTG GAGGTGCCTT ACTCAGTCTG AATCCAGGGA	1920
CCCTGAGCGG TGCTCTCAGC CCAGCTCTAA TGAGCAACAG TAACTGGCA ACTATTCAAG	1980
CTCTTGCTTC TGGTGGCTCT CTCCAATAA CATCACTTGA TGCAACTGGG AACCTGGTAT	2040
TTGCCAATGC GGGAGGAGCC CCCAACATCG TGAAGTCCCC TCTGTTCTTG AACCTCAGA	2100
ACCTCTCTCT GCTCACCAGC AACCTGTGTA GCTTGGTCTC TGCCGCCGCA GCATCTGCAG	2160
GGAACCTCTG ACCTGTAGCC AGCCTTCACG CCACCTCCAC CTCTGCTGAG TCCATCCAGA	2220
ACTCTCTCTT CACAGTGGCC TCTGCCAGCG GGGCTGCGTC CACCACCACC ACCGCCTCCA	2280
AGGCACAGTG AGCTGGGCAG AGCTGGGCTG CCAGAAGCCT TTTTCACTCT GCAGTGTGAT	2340

TGGACTGCCA GCCAGGTAA TAACTGAAA AATGTGATTG GCTTCCTCTC GCCGTGTTGT	2400
GAGGGCAAAG GAGAGAAGGG AGAAAAAAAA AAAAACC ACACACACCC ATACACAATA	2460
TACCAGAAAA GGAAGGAAGG ATGGAGACGG AACATTTGCC TAATTTGTAA TAAACACTG	2520
TCTTTTCAGG GTTGCTTCAT GGGTTGGAGG ACTTTCTAAC CAAAAATTAA AAAAAAAAAA	2580
AAAA	2584